

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 0 973 217 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
19.01.2000 Bulletin 2000/03

(51) Int. Cl.⁷: H01M 4/48, H01M 4/50,
H01M 4/52, H01M 4/04

(21) Application number: 99113425.5

(22) Date of filing: 12.07.1999

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 13.07.1998 JP 19785398
20.11.1998 JP 33144998

(71) Applicant: NGK INSULATORS, LTD.
Nagoya-City, Aichi Prefecture 467-8530 (JP)

(72) Inventors:
• Nemoto, Hiroshi
Nagoya-city, Aichi-pref. 457-0022 (JP)
• Takahashi, Michio
Nagoya-city, Aichi-pref. 467-0059 (JP)
• Kitoh, Kenshin
Nagoya-city, Aichi-pref. 458-0827 (JP)

(74) Representative:
Bühling, Gerhard, Dipl.-Chem. et al
Patentanwaltsbüro
Tiedtke-Bühling-Kinne & Partner
Bavariaring 4
80336 München (DE)

(54) Lithium transition metal compound for lithium secondary battery

(57) A lithium secondary battery has small internal resistance and has good charge-discharge cycle characteristics, with a lithium transition metal compound being used as a positive active material. A portion of transition element Me in a lithium transition metal compound LiMe_xO_y to be used as a positive active material is substituted by not less than two kinds selected from the group consisting of Li, Fe, Mn, Ni, Mg, Zn, B, Al, Co, Cr, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W to constitute $\text{LiM}_2\text{Me}_{x-z}\text{O}_y$. Here, M represents substitution elements, and $\text{M} \neq \text{Me}$.

EP 0 973 217 A2

Description

Background of the Invention and Related Art Statement

5 [0001] The present invention relates to, among secondary batteries to be used as a operational power source for a portable electric equipment, or as a motor driving battery for an electric vehicle or a hybrid electric vehicle, etc., a lithium secondary battery which has small internal resistance and has good charge-discharge cycle characteristics, with a lithium transition metal compound being used as a positive active material.

10 [0002] In recent years, miniaturization to go with lighter weight is being underway in an accelerated fashion on an electric equipment such as a personal handy phone system, a video tape recorder, a notebook-sized personal computer, etc., and a secondary battery comprising a lithium transition metal compound as a positive active material, with a carbon material as a negative material, and an electrolyte obtained by dissolving a Li ion electrolyte in an organic solvent, has become in common use as its power source battery.

15 [0003] Such a battery is generally called a lithium secondary battery or a lithium ion battery, and since they are provided with larger energy density as well as with higher unit cell voltage of approximate 4V, attention is being paid to it not only for the aforementioned handy electric equipment but also as a motor driving power source for an electric vehicle or a hybrid electric vehicle which is under consideration for a positive proliferation to the general public as a low pollution vehicle, on the background of recent environmental problems.

20 [0004] In such a lithium secondary battery, its battery capacity as well as its charge-discharge cycle characteristics (hereinafter called "cycle characteristics") heavily depends on material characteristics of a positive active material to be used. The lithium transition metal compound to be used as a positive active material includes lithium cobalt oxide ($LiCoO_2$), lithium nickel oxide ($LiNiO_2$), or lithium manganese oxide ($LiMn_2O_4$), etc. in particular.

25 [0005] Here, $LiCoO_2$ as well as $LiNiO_2$ comprises such features as a large Li capacity, a simple configuration, and excellent reversibility, and that it is provided with two dimensionally layered configuration being excellent in ion diffusion.

30 [0006] On the other hand, however, as concerns $LiCoO_2$, producing areas of Co are limited and it hardly is true that output quantity is abundant, and accordingly are expensive, thus there is a cost issue and there is a problem that its output density is smaller compared with $LiMn_2O_4$. In addition, as concerns $LiNiO_2$, synthesis of compound of stoichiometric composition is difficult since trivalent status of Ni is comparatively unstable, and in the case where detachment of Li becomes abundant, Ni will become subject to transition to bivalent status, emitting oxygen to constitute NiO , which creates problems such that a battery will stop functioning as a battery but also a risk of battery burst due to oxygen detachment may arise.

35 [0007] On the contrary, $LiMn_2O_4$ has a feature that raw materials are inexpensive and larger output density as well as higher voltage is provided. However, in the case where $LiMn_2O_4$ has been used as a positive active material, there is a problem that repetition of charging-discharging cycle gradually decreases discharge capacity and good cycle characteristics will not become obtainable. It is deemed that the major cause of this is reduction of the positive capacity since crystal configuration changes irreversibly due to insertion and detachment of Li^+ .

40 [0008] Thus, a lithium transition metal compound such as $LiCoO_2$, etc. respectively has both advantages and disadvantages together as a positive active material, and therefore, there are no rules which substances must be used, and it is deemed advisable that a positive active material which can show an appropriate feature for a particular purpose should be suitably picked and chosen for use.

45 [0009] Incidentally, regardless of the kind of a positive active material, it is preferred in terms of characteristics of a battery that the internal resistance of a battery is small, and it is a common problem to all the positive active materials to be solved that resistance in a positive active material (namely electronic conduction resistance) should be reduced, or in other words, electronic conductivity should be improved for this reduction of the internal resistance. Particularly, in a lithium secondary battery of large capacity used as a motor driving battery for an electric vehicle, etc., it is very important to obtain large current output necessary for acceleration and gradeability, etc. to improve charging-discharging efficiency.

50 [0010] Under the circumstances, conventionally, trials to improve electronic conductivity by adding to a positive active material conductive fine grains such as acetylene black, etc. to reduce internal resistance of a battery have been conducted. This is caused by that the above-described lithium transition metal compound is a mixed conducting body comprising both lithium ion conductivity and electronic conductivity together but its electronic conductivity is not always strong.

55 [0011] However, there is a problem that addition of acetylene black causes reduction of filling quantity of a positive active material to reduce battery capacity. In addition, it is deemed that improvement of electronic conductivity is not unlimited since acetylene black is a kind of carbon and is a semiconductor. Moreover, acetylene black is voluminous and presents such a problem that it is difficult to handle when an electrode plate is to be produced. Accordingly, the volume of its addition is to be limited to an appropriate quantity, comparing and considering advantageous effect of reduction of internal resistance, disadvantageous effect of reduction of battery capacity, and simplicity in production, etc.

[0011] Now, as described above, in the case where acetylene black has been added, acetylene black exists only on surfaces of particles of a positive active material, resulting in contributing to improvement of electronic conductivity among particles of positive active material, but not resulting in contributing to improvement of electronic conductivity inside a particle of a positive active material. Thus, conventionally, for improving electronic conductivity of a positive active material, attention was only paid to electronic conductivity among particles of a positive active material, but relationship between diffusion of Li^+ and electronic conductivity inside a particle of a positive active material at the time of battery reaction was not regarded as a problem.

[0012] In short, detachment of Li^+ from a particle of a positive active material as well as insertion of Li^+ to a particle of a positive active material is proceeded by diffusion of Li^+ inside a particle of a positive active material, simultaneously accompanied by transfer of electrons taking place inside a particle of a positive active material, and at this time, if electronic conductivity inside a particle of a positive active material is low, diffusion of Li^+ hardly is apt to take place and velocity of detachment and insertion of Li^+ , namely velocity of battery reaction, becomes slow, resulting in increase in internal resistance, which was not taken into consideration at all.

[0013] The present inventors paid attention to this point, and considered in earnest to improve electronic conductivity of a positive active material itself so that diffusion of Li^+ inside a positive active material may be proceeded well, thus reducing resistance of the positive active material itself, and at the same time, when a battery has been assembled without increasing volume of acetylene black to be added, internal resistance of that battery may be reduced, and as a result the present invention has been achieved.

20 Summary of the Invention

[0014] According to an aspect of the present invention, there is provided a lithium secondary battery, comprising a lithium transition metal compound LiMe_xO_y , in which a portion of transition element Me is substituted by not less than two kinds selected from the group consisting of Li, Fe, Mn, Ni, Mg, Zn, B, Al, Co, Cr, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W to constitute $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ (herein M represents substitution elements, and $M \neq Me$, and Z represents quantity of substitution.), the $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ being to be used as a positive active material.

[0015] In the present invention, not less than 2 kinds of elements are preferably selected as the substitution elements M among the above-described group of elements, particularly Li, Fe, Mn, Ni, Mg, Zn, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W, and it is especially preferred that at least Ti is included. It is also preferred that a portion of the remaining transition elements Me in $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ to include not less than two kinds of substitution elements M to be obtained this way is also preferably substituted further by at least not less than one kind of elements selected among B, Al, Co, and Cr. Also it is preferred that in a lithium transition metal compound $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$, Z/X , the ratio of the substitution quantity Z of substitution elements M and Me quantity X of the original transition elements, fulfills the condition of $0.005 \leq Z/X \leq 0.3$.

[0016] Incidentally, as one of lithium transition metal compound to be suitably used in the present invention, lithium manganese oxide, especially a lithium manganese oxide having a spinel configuration of cubic system, may be nominated. The average valence of substitution elements M to substitute a portion of manganese in such lithium manganese oxide is set at not less than 3 but not more than 4. Here, an average valence is an average value of ion valence of not less than two different substitution elements M in a positive active material. Here, in the case where lithium manganese oxide has been used, a substitution quantity Z preferably remains within a range of $0.01 \leq Z \leq 0.5$ and more preferably fulfills a condition of $0.1 \leq Z \leq 0.3$.

[0017] In addition, in the present invention, lithium cobalt oxide or lithium nickel oxide is suitably used as a lithium transition metal compound. In the case where such materials have been used, it is preferred that the average valence of substitution elements M to be substituted with a portion of cobalt or nickel in lithium cobalt oxide or lithium nickel oxide is 3. However, the case where all the substitution elements M have the ion valence of 3 is excluded. Here, the substitution quantity Z preferably remains within the range of $0.005 \leq Z \leq 0.3$, and further preferably fulfills the condition of $0.05 \leq Z \leq 0.3$.

[0018] $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ to be used in the above-described lithium secondary battery of the present invention is composed by firing a mixed compound comprising salts and/or oxides having been prepared with a predetermined ratio in oxidation atmosphere at a range of 600°C to 1000°C , spending 5 hours to 50 hours. At this time, also suitably adopted is such a method that is conducted, dividing firing into not less than twice, with the firing temperature for the forthcoming step to be set higher than that for the previous step, and thus proceeding with composition. Here, in the case where plurality of firing is conducted, the final firing is to be conducted under a firing condition involving oxidation atmosphere at a range of 600°C to 1000°C , spending 5 hours to 50 hours.

55

Detailed Description of Preferred Embodiment

[0019] In a lithium secondary battery of the present invention, a portion of transition element Me of a lithium transition

metal compound LiMe_xO_y is substituted by not less than two kinds of elements to constitute $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$, the $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ being to be used as a positive active material. Here, M represents substitution elements, and substitution elements M are the one which are different from a transition element Me ($M \neq \text{Me}$), and Z represents quantity of substitution. Strictly, since not less than two kinds of substitution elements M are involved, the chemical formula of the positive active material is described as $\text{Li}((\text{M}_1)^{x_1}(\text{M}_2)^{x_2} \dots (\text{M}_n)^{x_n})_z\text{Me}_{x-z}\text{O}_y$ (herein, $\text{M}_1, \text{M}_2, \dots, \text{M}_n$ represent respectively different elements, and the total sum of x_1 to x_n is 1) for substitution by n-numbered kinds of elements. Incidentally, element substitution of the present invention involving such plural elements will be hereinafter called "complex substitution".

[0020] As substitution elements M, not less than two kinds of elements are selected from the group consisting of Li, Fe, Mn, Ni, Mg, Zn, B, Al, Co, Cr, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W. These elements were determined by applying Hume-Rothery's rule to an ionic radius introduced by SHANNON, et al which has been described in *Acta Cryst.* (1976) A32, 751, and for the ion radius of transition element Me to be substituted in a space group $R(-3)m$ (herein "-" represents rotation-inversion) or in $Fd\bar{3}m$ (a spinel configuration), a condition that the coordination number for oxygen is the same as that for the transition element Me and the average ionic radius of the substitution elements M remains within $\pm 15\%$ of the ionic radius of the transition element Me, and is not a radioactive element nor a gas, and not strongly toxic having been fulfilled so as to select a combination of elements. Here, as a transition element Me, Mn, Co, and Ni to be suitably used in the present invention are regarded as a standard.

[0021] An ionic radius of substitution elements M is referred to an average value of ionic radius of not less than 2 kinds of elements, and is determined in consideration of existence ratio of each element. In the present invention, it is preferable that all the ionic radius of the substitution elements M remains within $\pm 15\%$ of the ionic radius of the transition element Me, but in the case where such a condition may not be fulfilled, for example even in the case of the substitution elements M_1 numbered 1 with its ionic radius far bigger outside the range of $+15\%$ of the ionic radius of the transition element Me, and the substitution elements M_2 numbered 2 with its ionic radius far smaller outside the range of -15% of the ionic radius of the transition element Me, if an average ionic radius of the substitution elements M_1 and M_2 falls in the range of $\pm 15\%$ of the ionic radius of the transition element Me, complex substitution is feasible.

[0022] However, in the case where Li is used, Li can be used as a substitution elements M, exceptionally, even when the above-described conditions on ionic radius are not fulfilled. The reasons of this are that other than the ionic radius of the above-described version of SHANNON, et al, there is also a version of Polling, et al, and there is a big difference in normal values for these versions, thus limiting consideration on only the ionic radius of Li is problematic in terms of character itself, and that Li is an original constitutional element and particularly in the LiMn_2O_4 system, Li is deemed to substitute the position of Mn, and further that it is experimentally possible to solid-solubilize Li.

[0023] Incidentally, as concerns substitution elements M, in theory, Li is to become +1 valence ion, Fe, Mn, and Ni, Mg, and Zn are +2 valence ions, B, Al, Co, and Cr are +3 valence ions, Si, Ti, and Sn are +4 valence ions, P, V, Sb, Nb, and Ta are +5 valence ions, and Mo and W are +6 valence ions, and they all are elements to be solid-solubilized in $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$. However, for Co and Sn, they can be +2 valence ions, and for Fe, Sb and Ti, they can be +3 valence ions and for Mn they can be +3 and +4 valence ions, and for Cr they can even be +4 and +6 valence ions.

[0024] Therefore, as seen in an actual positive active material, in the case where there exists a part of ionic valence subject to change in valence values due to various crystallographic deficiency, in some cases there is a possibility that an average valence of substitution elements M might not coincide with theoretic valence value, e.g. 3.5 for lithium manganese oxide and 3 for lithium cobalt oxide as well as lithium nickel oxide, of a transition element Me prior to complex substitution.

[0025] For example, since Ti can exist comparatively stably under +3 valence condition in addition to +4 valence condition, in the case where Ti has been solid-solubilized in $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ under the condition having such mixed atomic valence, the average valence of Ti falls in a range between +3 to +4. And as concerns Fe, since Fe remains equally stable under +2 and +3 valence condition and it is also known that the status of +4 valence exists stably in a certain chemical compound, the average valence of Fe in $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ is to fall in a range between +2 to +4. In addition, similarly, also as concerns quantity of oxygen in $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$, it may exist in deficit or in excess within a range to sustain a crystal configuration.

[0026] Incidentally, as a lithium transition metal compound to be used in the present invention, lithium manganese oxide, lithium cobalt oxide, and lithium nickel oxide may be nominated in particular. Here as concerns lithium manganese oxide, a lithium manganese oxide (LiMn_2O_4) having a spinel configuration of cubic system is suitably used. In LiMn_2O_4 , one Mn in two units of Mn is in the state of +3 valence while the other Mn is in the state of +4 valence state. Accordingly, in complex substitution, two cases can be considered, namely a case where substitution elements M is used for substitution of Mn in this +3 valence state, and a case involving substitution of Mn in +4 valence state.

[0027] An average valence value of the substitution elements M is 3 in the case where complex substitution of +3 valence Mn takes place, but here at Least elements to become ions with other than +3 valence is included in the substitution elements M. For example, such cases that two units of +3 valence Mn undergo complex substitution with one +2 valence Mg and +4 valence Ti, and two units of +3 valence Mn undergo substitution with one +1 valence Li and one +5 valence V can be nominated. And in the case where a +3 valence Mn undergoes complex substitution with such an

element having other than +3 valence, it is permitted that the remaining +3 valence Mn is substituted with another +3 valence ion. Here, an average valence is referred to an average value of ion valence of not less than two different substitution elements M in a positive active material and is determined, putting their existence ratio under consideration.

[0028] Likewise, for the purpose that +4 valence Mn undergoes complex substitution, it is necessary that substitution has taken place with at least an element to provide a valence value other than +4 valence, and thereafter the remaining +4 valence Mn may be substituted with an element to provide the same +4 valence. In general, in complex substitution of LiMn_2O_4 , at least it is necessary that the ionic valence of the substitution elements M numbered 1 is not more than 3 and the ionic valence of another substitution elements M is not less than 4, consequently resulting in the average valence of only substitution elements M to be ranged from not less than 3 to not more than 4, and the average valence value obtained from the substitution elements M after complex substitution inclusive of Mn being 3.5.

[0029] On the other hand, since the substitution elements M to make a portion of Co or Ni in lithium cobalt oxide (LiCoO_2), and lithium nickel oxide (LiNiO_2) undergo complex substitution is to provide an average valence value of 3, similarly in the above-described substitution of +3 valence Mn, the substitution elements M are to include elements to provide ions with at least other than +3 valence. Therefore, the case where all the substitution elements M have ionic valence value of 3 valence is excluded from complex substitution of the present invention.

[0030] In the case where a battery has been assembled using a positive active material which had undergone such complex substitution, there reveals an effect with remarkable reduction of internal resistance. This is deemed to be caused by that electronic conductivity is improved in the frame of lithium transition elemental composite compound (a portion exclusive of Li attributable to ionic conduction), and thus velocity of detachment and insertion of Li ions in battery reaction has become faster. And considering that the lattice constant gets small due to complex substitution, the improvement of electronic conductivity in this frame is presumed to heavily depend on that in the case where transition elements Me each other and/or substitution elements M are transition metal elements, the d orbital between substitution elements M and a transition element Me is apt to overlap, which makes it easier to smoothly proceed with electrons' movement by use of this d orbital.

[0031] In addition, repeating charge and discharge of a battery assembled by use of materials which have undergone complex substitution, no deterioration is observed, compared with the case involving use of materials which have not undergone complex substitution, and therefore, it is deemed that complex substitution does not negatively affect stability of the frame. Moreover, in LiMn_2O_4 , as shown in the below-described embodiments, the cycle characteristics have been improved, thus it is deemed that complex substitution attributes to improvement of reversibility of crystal lattice associated with insertion and detachment of Li ions.

[0032] Incidentally, compared with the case where a portion of the transition element Me is substituted by another element (hereinafter, such substitution involving one element is referred to as "single element substitution."), according to complex substitution, such a problem that positive capacity might be reduced by larger volume of substitution in single element substitution can be avoided. Next, this example is explained by use of LiMn_2O_4 , but it goes without saying that the explanation may be made to LiCoO_2 and LiNiO_2 .

[0033] In the case where Mn^{3+} in LiMn_2O_4 has undergone single element substitution with an element having valence value of not more than two valence, e.g. one valence ion such as Li^+ , charge equivalent to +2 valence value, being a difference of charge with Mn^{3+} , will be in short, thus for the purpose of maintaining electrical neutrality of materials, two units of Mn^{3+} will be changed to Mn^{4+} . Thus, consequently, one Li^+ will be substituted with Mn^{3+} and solid-solubilized, resulting in reduction of approximately three units of Mn^{3+} .

[0034] Here, in LiMn_2O_4 , it is deemed that, at the time of charging, electrical neutrality of materials is maintained by compensating shortage of charge due to detachment of Li^+ with Mn^{3+} being changed to Mn^{4+} , and at the time of discharging reverse reaction takes place. In short, the quantity of Mn^{3+} in LiMn_2O_4 determines the positive capacity, and a quantity of Li^+ corresponding to Mn^{3+} attributes charging and discharging reaction. Therefore, for the purpose that Li^+ is detached from a crystal lattice or inserted into a crystal lattice, it will become necessary that a change in valence value takes place in cations other than Li^+ , namely substitution elements M and/or transition element Me.

[0035] However, in the previous embodiment, Li^+ which was substituted with Mn^{3+} has not undergone change in valence value, consequently Mn^{3+} remains in short by three units. Therefore, 3 units of Li^+ will not attribute to charging and discharging reaction. In short, consequently there arises a problem that the positive capacity is reduced in excess of quantity of substitution. Such a problem similarly takes place in single element substitution involving +2 valence ions.

[0036] On the other hand, in complex substitution of the present invention, substitution elements M are to be narrowed to Li, Fe, Mn, Ni, Mg, Zn, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W (hereinafter these substitution elements M are referred to as "substitution elements group within a reduced range."), and at least not less than two kinds of elements are arranged to be selected, then in addition to an effect that improves electronic conductivity, the above-described problem that the positive capacity is reduced in excess of quantity of element substitution is avoided.

[0037] In short, when ions with +1 valence or +2 valence and ions with +4 to +6 valence are combined, as concerns shortage of positive charge caused by solid-solubilizing ions with +1 valence or +2 valence, the charge is not compensated by change of Mn^{3+} to Mn^{4+} , but ions with +4 to +6 valence are solid-solubilized and compensated, thus without

reducing the positive capacity as a result of reducing the number of Mn^{3+} in excess of substitution quantity, Mn can undergo substitution.

[0038] For example, in the case where two units of Mn^{3+} are substituted by one Li^+ and one V^{5+} , reduction of positive capacity is limited to a reduced volume of two units of Mn^{3+} , and it will become possible to make quantity of reduction of Mn^{3+} lesser than reduction by three units of Mn^{3+} in the case where one Mn^{3+} has undergone single element substitution with one Li^+ . In addition, in the case where two units of Mn^{3+} has been substituted with one Mg^{2+} and one T^{4+} , reduction of positive capacity is limited to reduction covering two units of Mn^{3+} , and is lesser than reduction of four units of Mn^{3+} in the case where two units of Mn^{3+} have been substituted with two units of Mg^{2+} . Thus, reduction quantity of Mn^{3+} is equivalent to substitution quantity of elements, and accordingly such event that reduction in positive capacity exceeds substitution quantity is to be avoided.

[0039] Here, in complex substitution, when at least Ti is arranged to be included as substitution elements M, a remarkable effect of improvement on electronic conductivity is obtainable and preferable. In addition, Ti can be effectively used to prevent a drop in positive capacity, which is preferable.

[0040] In $LiM_2Me_{X-Z}O_Y$ including not less than two kinds of substitution elements M obtainable when complex substitution using elements among the above-described substitution elements group within a reduced range, a portion of remaining transition elements Me may further be substituted with at least not less than one kind of element selected from B, Al, Co, and Cr. In this case, complex substitution involving at least three kinds of element is to take place.

[0041] These elements such as B and Al, etc. exist in $LiM_2Me_{X-Z}O_Y$ as ions with +3 valence in theory. But, as described above, in actual positive active materials, the ion valence value does not always have to correspond with the theoretic valence values. Ions with +3 valence is substituted with Mn^{3+} one on one, therefore, decrease in positive capacity is the same as the quantity of substitution, and decrease in positive capacity not less than the quantity of substitution does not take place, and on the other hand, the said ion attributes to improvement of electron conductivity of a positive active material itself. Incidentally, in the case where $LiMn_2O_4$ is used, an effect that its crystal configuration is made reversible toward insertion and detachment of Li^+ is provided.

[0042] Next, substitution quantity Z in complex substitution is explained. In the present invention, it is preferred that Z/X , the ratio of the quantity Z to be substituted by substitution elements M to the quantity X of the original transition element Me fulfills the condition of $0.005 \leq Z/X \leq 0.3$. When Z/X is less than 0.005, resistance of a positive active material does not drop, and improvement in cycle characteristics rarely appears. In short, no effects of complex substitution appear. On the other hand, when Z/X is more than 0.3, in synthesis of a positive active material, production of a different phase is admitted through powder X-ray diffraction method (XRD), and a single phase material was not obtainable. In a battery, such a different phase only increases the weight of a positive active material and does not attribute to battery reaction, thus it goes without saying that production of a different phase at the time of synthesis together with entry to the battery should be avoided.

[0043] Positive-active-material-wise, in particular, when $LiMn_2O_4$ has been used, the substitution quantity Z is preferably to fall in within a range of $0.01 \leq Z \leq 0.5$, and further preferably to fall in a range of $0.1 \leq Z \leq 0.3$, and when $LiCoO_2$ as well as $LiNiO_2$ is used, the substitution quantity Z is preferably to fall in within a range of $0.005 \leq Z \leq 0.3$, and further preferably to fall in a range of $0.05 \leq Z \leq 0.3$, and within the respective preferable ranges of the substitution quantity Z, there remarkably appears an effect of improvement of electronic conductivity of a positive active material, which is preferable.

[0044] Incidentally, when elemental substitution by not less than one kind to have been selected from B, Al, Co, and Cr further took place as well after complex substitution, the total substitution quantity ($Z+w$) of substitution quantity Z of substitution elements M to have been selected from a group of substitution elements within a reduced range, and the substitution quantity (to be referred to as "w") of B and Al, etc. is required to fulfill a relationship of $0.01 \leq Z+w \leq 0.5$.

[0045] Incidentally, $LiM_2Me_{X-Z}O_Y$ to be used in a lithium secondary battery of the present invention, is composed by firing a mixed compound comprising salts and/or oxides of each element (substitution elements M as well as Li and transition element Me) having been prepared with a predetermined ratio in oxidation atmosphere at a range of 600°C to 1000°C, spending 5 hours to 50 hours, and thus a single-phase product can be obtained. Here, an oxidation atmosphere is referred to as an atmosphere having partial pressure of oxygen with which generally a sample inside a furnace is brought into oxidation reaction. In synthesis of $LiCoO_2$ as well as $LiNiO_2$, it is preferable that the partial pressure of oxygen is set at not less than 10 %, and, in particular, air atmosphere and oxygen atmosphere, etc. are applicable.

[0046] Incidentally, when the firing temperature is as low as less than 600°C, a peak showing residue of raw material, e.g. peak of lithium carbonate (Li_2CO_3) in the case where Li_2CO_3 is used as a lithium source, is to be observed in XRD chart of fired product, and no single phase products can be obtained. On the other hand, when the firing temperature is as high as more than 1000°C, high temperature phase is produced in other than a compound of the targeted crystal system, and a single phase product will become no longer obtainable.

[0047] In addition, firing may be conducted, being divided into not less than twice. In that case, it is preferable that the firing is proceeded with the firing temperature for the forthcoming step to be set higher than that for the previous step, and the final firing is to be conducted under a firing condition involving oxidation atmosphere at a range of 600°C to

1000°C, spending 5 hours to 50 hours. Thus, in the case of firing taking place twice for example, applying this condition of second firing temperature as well as firing period, the product obtainable when synthesis has been conducted with the temperature for the second firing to be set at not less than the temperature for the first firing features steeper projection in the peak shape in the XRD chart than with the product obtainable when a single firing yields, and as a result improvement of crystallinity can be planned.

5 [0048] A salt for each element will not be limited in particular, but it goes without saying that those having intensive purity and further being inexpensive as raw materials are preferably to be used. Accordingly, such carbonate, hydroxide, and organic acid/salt that do not produce harmful decomposition gas at the times of elevation of temperature or firing are preferably used. However, nitrate, hydrochloride, and sulfate, etc. are not always unusable. Generally, in synthesis 10 of LiCoO_2 and LiNiO_2 , it is known that synthesis temperature goes down with usage of salts instead of oxides as raw materials. Here, as concerns raw materials on Li, usually an oxide Li_2O is chemically unstable, and thus it is rarely used.

15 [0049] As the foregoing, implementation of complex substitution of the present invention will make improvement in electronic conductivity of a positive active material easier to plan, providing preferable electric characteristics, and resulting in decrease in internal resistance of a battery. In addition, the problem that positive capacity might be reduced by larger volume of element substitution in single element substitution which conventionally used to be problematic in single element substitution, is to be solved, and reduction of positive capacity is to be suppressed to the extent equivalent to quantity of element substitution. At the same time, as concerns LiMn_2O_4 , reversibility of crystal configuration for insertion and detachment of Li^+ is improved, thus cycle characteristics as a battery is improved. Accordingly, decrease with the passage of time in battery capacity due to repetition of charging and discharging is controlled.

20 [0050] Reduction of internal resistance and reservation of positive capacity, and increase in cycle characteristics are planned in such a battery, which is used as a motor driving power source for an EV or an HEV in particular, consequently providing with an excellent effect that predetermined running performance such as acceleration performance as well as slope-climbing performance, etc. is maintained, and continuous running distance per a charging is kept for long.

25 [0051] Incidentally, other materials to be used for production of a battery are not limited to whatsoever, and conventionally known various materials can be used. For example, as a negative active material, an amorphous carbon material such as soft carbon or hard carbon, or carbon material such as artificial graphite such as high graphitized carbon material, etc. and natural graphite, etc. is used.

30 [0052] And as an organic electrolyte a carbonic acid ester family such as ethylene carbonate (EC), diethyle carbonate (DEC), and dimethyle carbonate (DMC), and the one in which one or more kinds of lithium fluoride complex compound such as LiPF_6 , and LiBF_4 , etc. or lithium halide such as LiClO_4 as an electrolyte are dissolved in a single solvent or mixed solvent of organic solvents such as propylene carbonate (PC), γ -butyrolactone, tetrahydrofuran, and acetonitrile, etc., can be used.

35 Example

40 [0053] Successively, taking as a major embodiment complex substitution involving two kinds of elements as substitution elements M including Ti which provides most remarkable effect in the present invention, based on whose experimental results an explanation is provided as follows:

(Synthesis of positive active material $\text{LiM}_{2-x}\text{Mn}_{2-x}\text{O}_4$)

45 [0054] As the starting raw material, powder of commercially available Li_2CO_3 , MnO_2 , TiO_2 , MgO , and NiO was used and was weighed and mixed so that the positive active material composition of respective embodiments shown in Table 1 (positive active materials for measurement of internal resistance ratio), Table 2 (positive active materials for measurement of capacity of the initial charging), and Table 3 (positive active materials for a cycle test) might be obtained, and firing took place at 800°C in the air atmosphere for 24 hours, and the positive active materials were obtained. Here, when the combination of substitution elements M took place involving Ti and Mg or Ni, the mixing ratio of them was set at $\text{Ti:Mg or Ni} = 1:1$, and for the case involving Li and Ti, it was set at $\text{Li:Ti} = 1:2$. Incidentally, for the purpose of comparing the effects of complex substitution and single element substitution, positive active materials where a portion of Mn underwent single element substitution with Mg, Ti, Ni, and Li respectively as well as LiMn_2O_4 which did not undergo element substitution were formed under the similar conditions.

[Table 1]

	Positive active material composition	Internal resistance ratio of coin cells (%)
Comparative example 1	LiMn ₂ O ₄	100
Embodiment 1	Li(Mg _{0.5} Ti _{0.5}) _{0.01} Mn _{1.99} O ₄	54
Embodiment 2	Li(Mg _{0.5} Ti _{0.5}) _{0.10} Mn _{1.90} O ₄	37
Embodiment 3	Li(Mg _{0.5} Ti _{0.5}) _{0.15} Mn _{1.85} O ₄	35
Embodiment 4	Li(Mg _{0.5} Ti _{0.5}) _{0.30} Mn _{1.70} O ₄	29
Embodiment 5	Li(Mg _{0.5} Ti _{0.5}) _{0.50} Mn _{1.50} O ₄	41
Embodiment 6	Li(Ni _{0.5} Ti _{0.5}) _{0.01} Mn _{1.99} O ₄	52
Embodiment 7	Li(Ni _{0.5} Ti _{0.5}) _{0.10} Mn _{1.90} O ₄	36
Embodiment 8	Li(Ni _{0.5} Ti _{0.5}) _{0.15} Mn _{1.85} O ₄	36
Embodiment 9	Li(Ni _{0.5} Ti _{0.5}) _{0.30} Mn _{1.70} O ₄	30
Embodiment 10	Li(Ni _{0.5} Ti _{0.5}) _{0.50} Mn _{1.50} O ₄	45
Comparative example 2	LiMg _{0.15} Mn _{1.85} O ₄	80
Comparative example 3	LiTi _{0.15} Mn _{1.85} O ₄	69
Comparative example 4	LiNi _{0.15} Mn _{1.85} O ₄	71

[Table 2]

	Positive active material composition	Capacity of the initial charging (mAh/g)
Embodiment 3	Li(Mg _{0.5} Ti _{0.5}) _{0.15} Mn _{1.85} O ₄	102
Embodiment 12	Li(Li _{0.33} Ti _{0.67}) _{0.15} Mn _{1.85} O ₄	102
Comparative example 2	LiMg _{0.15} Mn _{1.85} O ₄	85
Comparative example 3	LiTi _{0.15} Mn _{0.85} O ₄	105
Comparative example 5	LiLi _{0.15} Mn _{1.85} O ₄	70

[Table 3]

	Positive active material composition	Capacity ratio toward capacity of the initial charging of a battery after 100 cycles
Embodiment 2	Li(Mg _{0.5} Ti _{0.5}) _{0.10} Mn _{1.90} O ₄	0.69
Embodiment 3	Li(Mg _{0.5} Ti _{0.5}) _{0.15} Mn _{1.85} O ₄	0.84
Embodiment 4	Li(Mg _{0.5} Ti _{0.5}) _{0.30} Mn _{1.70} O ₄	0.83
Embodiment 5	Li(Mg _{0.5} Ti _{0.5}) _{0.50} Mn _{1.50} O ₄	0.73
Embodiment 11	Li(Li _{0.33} Ti _{0.67}) _{0.10} Mn _{1.90} O ₄	0.71
Embodiment 12	Li(Li _{0.33} Ti _{0.67}) _{0.15} Mn _{1.85} O ₄	0.85

[Table 3] (continued)

	Positive active material composition	Capacity ratio toward capacity of the initial charging of a battery after 100 cycles	
5	Embodiment 13	Li(Li _{0.33} Ti _{0.67}) _{0.30} Mn _{1.70} O ₄	0.82
10	Embodiment 14	Li(Li _{0.33} Ti _{0.67}) _{0.50} Mn _{1.50} O ₄	0.70
15	Comparative example 2	LiMg _{0.15} Mn _{0.85} O ₄	0.68
20	Comparative example 3	LiTi _{0.15} Mn _{1.85} O ₄	0.66
25	Comparative example 5	LiLi _{0.15} Mn _{1.85} O ₄	0.69

(Synthesis of positive active materials LiM_zCo_{1-z}O₂ and LiM_zNi_{1-z}O₂)

[0055] As the starting raw material, commercially available Li₂CO₃, Co₃O₄, NiO, MgO, and TiO₂ were used and were weighed and mixed so that the composition of respective kinds of embodiments shown in Table 4 as well as Table 5 (positive active materials for measurement of internal resistance ratio) might be obtained. And as concerns LiM_zCo_{1-z}O₂, firing took place at 900°C in the air atmosphere for 20 hours, and on the other hand as concerns LiM_zNi_{1-z}O₂, firing took place at 750°C in an oxygen atmosphere for 20 hours to proceed with synthesis. In addition, as put down in Table 4 as well as Table 5, LiCoO₂ as well as LiNiO₂ in which no addition elements were added, and also samples related to Examples undergoing single element substitution were synthesized under the similar conditions. The formed respective kinds of positive active materials of Embodiments as well as Comparative examples were confirmed to be in a single phase through XRD.

[Table 4]

	Positive active material composition	Internal resistance ratio of coin cells (%)	
30	Comparative example 6	LiCoO ₂	100
35	Embodiment 15	Li(Mg _{0.5} Ti _{0.5}) _{0.005} Co _{0.995} O ₂	86
40	Embodiment 16	Li(Mg _{0.5} Ti _{0.5}) _{0.05} Co _{0.95} O ₂	69
45	Embodiment 17	Li(Mg _{0.5} Ti _{0.5}) _{0.25} Co _{0.75} O ₂	65
50	Embodiment 18	Li(Mg _{0.5} Ti _{0.5}) _{0.3} Co _{0.7} O ₂	73
	Embodiment 19	Li(Ni _{0.5} Ti _{0.5}) _{0.005} Co _{0.995} O ₂	88
	Embodiment 20	Li(Ni _{0.5} Ti _{0.5}) _{0.05} Co _{0.95} O ₂	63
	Embodiment 21	Li(Ni _{0.5} Ti _{0.5}) _{0.25} Co _{0.75} O ₂	59
	Embodiment 22	Li(Ni _{0.5} Ti _{0.5}) _{0.3} Co _{0.7} O ₂	67
	Comparative example 7	LiMg _{0.05} Co _{0.95} O ₂	90
	Comparative example 8	LiTi _{0.05} Co _{0.95} O ₂	87
	Comparative example 9	LiNi _{0.05} Co _{0.95} O ₂	94

[Table 5]

	Positive active material composition	Internal resistance ratio of coin cells (%)	
55	Comparative example 10	LiNiO ₂	100
	Embodiment 23	Li(Li _{0.33} Ti _{0.67}) _{0.005} Ni _{0.995} O ₂	91

[Table 5] (continued)

	Positive active material composition	Internal resistance ratio of coin cells (%)	
5	Embodiment 24	Li(Li _{0.33} Ti _{0.67}) _{0.05} Ni _{0.95} O ₂	77
	Embodiment 25	Li(Li _{0.33} Ti _{0.67}) _{0.25} Ni _{0.75} O ₂	72
10	Embodiment 26	Li(Li _{0.33} Ti _{0.67}) _{0.3} Ni _{0.7} O ₂	80
	Comparative example 11	LiTi _{0.05} Ni _{0.95} O ₂	93
	Comparative example 12	LiLi _{0.05} Ni _{0.85} O ₂	110

(Forming of a battery)

15 [0056] At first, using the formed various kinds of positive active materials, and mixing a positive active materials, acetylene black powder being conductive material, and polyvinylidene fluoride being bonding material with a weight ratio of 50:2:3 to form a positive material. A disk shape having diameter of 20 mmØ was prepared as a positive pole by press-forming 0.02 g of the said positive material under a pressure of 300 kg/cm². Next, in accordance with test purposes, as described below, two kinds of coin cells were formed. In short, the coin cells for measuring internal resistance set forth
20 in Table 1, Table 4, and Table 5 as well as the coin cells for cycle tests set forth in Table 3 were formed by using a positive pole having been formed as described above, an electrolyte having been formed by dissolving LiPF₆ being an electrolyte into an organic solvent with ethylene carbonate and diethyle carbonate being mixed with a same volume ratio to constitute a density of 1 mol/L, a negative pole made of carbon, and a separator separating the positive electrode and the negative pole.

25 [0057] On the other hand, the coin cells for measuring the capacity for initial charging set forth in Table 2 was formed by using a positive pole having been formed, an electrolyte having been formed by dissolving LiClO₄ being an electrolyte into propylene carbonate to constitute a density of 1 mol/L, a negative pole made of metal Li, and a separator separating the positive pole and the negative pole.

30 (Method to measure a battery's internal resistance and the results thereof)

35 [0058] As concerns coin cells having been formed as described above, using respective kinds of positive active materials set forth in Table 1, Table 4 and Table 5, only one cycle of charging and discharging test was conducted, involving charging constant current of 1C rate and constant voltage of 4.1 V in accordance with the capacity of a positive active material, and similarly discharging constant current of 1C rate and constant voltage 2.5 V at a, and the battery's internal resistance was obtained by dividing difference between the potential at a resting state after finishing charging and the potential immediately after commencement of discharging (potential difference) with discharging currency. And the internal resistance of a battery using a positive active material which underwent single element substitution and complex substitution was divided by the internal resistance of a battery using original compound which did not undergo elemental substitution respectively (LiMn₂O₄, LiCoO₂, and LiNiO₂) to yield a value, which was stipulated as an internal resistance ratio. Accordingly, as the value of internal resistance ratio gets smaller, reduction effect on internal resistance gets larger. The results have been put down in Table 1, Table 4, and Table 5 respectively.

45 [0059] Based on Table 1, as concerns LiMn₂O₄, in the case where positive active materials having undergone single element substitution were used, in other words, in the case where the embodiments 1 through 10 having involved positive active materials having undergone complex substitution while internal resistance in comparative examples 2 through 4 has halted at approximately 70 % at the best, it is obvious that the substitution quantity Z has fallen in the range of 0.01 ≤ Z ≤ 0.5, and the internal resistance ratio has been decreased to reach not more than approximately 50 %. In addition, as shown in embodiments 2 through 4 as well as embodiments 7 through 9, in the case where complex substitution has taken place so that substitution quantity Z may fall in the range of 0.1 ≤ Z ≤ 0.3, it is obvious that remarkable reduction effect in internal resistance has been obtained.

50 [0060] Based on Table 4, as concerns LiCoO₂, compared with comparative examples 7 through 9 where single element substitution took place, it was confirmed that remarkable reduction in internal resistance appeared in embodiments 15 through 22 where complex substitution took place. And, as shown in embodiments 16 through 18 as well as embodiments 20 through 22, for the range of 0.1 ≤ Z ≤ 0.3, Z being substitution quantity, remarkable reduction effect in internal resistance has appeared. Incidentally, in the case where LiCoO₂ is the basic material, reduction effect in internal resistance has been limited to a small extent, compared with the case involving LiMn₂O₄.

55 [0061] The value of internal resistance ratio obtained by single element substitution as well as complex substitution having used LiNiO₂ as the basic material has been similar to the case having involved LiCoO₂, and compared with com-

parative examples 11 and 12 where single substitution took place, the internal resistance ratio has been reduced to a large extent in embodiments 23 through 26 where complex substitution took place, and as shown in embodiments 24 through 26, for the range of $0.05 \leq Z \leq 0.3$, Z being substitution quantity, a reduction effect in internal resistance has appeared to a large extent. However, as in the case of LiCoO_2 , compared with the case using LiMn_2O_4 , the effect of decrease in internal resistance is little.

[0062] From these results, complex substitution by not less than two kinds selected from the group consisting of Li, Fe, Cr, Mn, Ni, Mg, Zn, B, Al, Co, Cr, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W has been conducted, and forming of positive active materials through measurement of internal resistance by the method similar to the one described above, and as a result the tendency similar to the case involving complex substitution having shown in Table 1 was confirmed.

(Measurement of a battery's internal charging capacity and the results thereof)

[0063] As concerns coin cells having been formed as previously described, using positive active materials set forth in Table 2, the initial charging capacity (battery capacity) was measured, involving charging to reach 4.2 V at a constant currency and constant voltage of 0.2C rate in accordance with the capacity of a positive active material. The results have been put down in Table 2. Based on these results, it is obvious that in the case where element substitution quantity as a whole is same, compared with single element substitution by Li^+ and Mg^{2+} respectively, battery capacity has got large in the case where complex substitution took place, however, in the case involving single element substitution by Ti^{4+} , battery capacity approximately equivalent to that in the case involving complex substitution has especially been obtained.

[0064] It is deemed that in single element substitution respectively by Li^+ and Mg^{2+} , as previously described, reduction in Mn^{3+} in the quantity not less than element substitution quantity has reduced Li^+ attributable to charging and discharging, and thus has reduced battery capacity, nevertheless, complex substitution has shown that it has controlled the said reduction in capacity. It is deemed on the other hand that in the case involving single element substitution by Ti^{4+} , most portion of Ti^{4+} has undergone change in valence value to Ti^{3+} at the time of firing, and thus, substitution between Ti^{3+} and Mn^{3+} have made available the battery capacity equivalent to that obtainable in complex substitution.

[0065] Having this result in hand, complex substitution by not less than two kinds selected from the group consisting of Li, Fe, Cr, Mn, Ni, Mg, Zn, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W has been conducted, and forming of positive active materials through assessment of battery capacity was conducted by the method similar to the one described above, and as a result the characteristics similar to the case involving complex substitution having shown in Table 2 were obtained.

[0066] In addition, for the purpose of looking into a range of composition where reduction controlling effect on battery capacity by complex substitution appears, experiments similar to those described above with variety of substitution quantity Z , it became obvious that the substitution quantity Z preferably fell in the range of $0.01 \leq Z \leq 0.5$. In the case where the substitution quantity Z exceeded 0.5, in any combination of substitution elements M, production of compounds other than those in the spinel phase was confirmed by XRD.

(Cycle operation test and the results thereof)

[0067] Successively, further for the purpose of looking into cycle characteristics in a substitution quantity Z where an effect of complex substitution reveals, as concerns batteries having been formed as previously described, using positive active materials having respective compositions set forth in Table 3, a cycle operation test was conducted, repeating charging constant current of 1C rate and constant voltage of 4.1 V and likewise discharging constant current of 1C rate and constant voltage of 2.5 V in accordance with the capacity of a positive active material.

[0068] In Table 3, discharging capacity of a battery after the consummation of 100 cycles has been put down in terms of ratio toward the initial discharging capacity of a battery. Consequently, as this ratio gets larger, reduction in battery's discharging capacity is deemed to get less. As shown in Table 3, it became obvious that in a battery where positive active materials having undergone complex substitution were used, reduction quantity in battery's discharging capacity is as a whole smaller than in the case involving positive active materials having undergone single element substitution, and the said reduction was little especially within a range of $0.1 \leq Z \leq 0.3$, Z being substitution quantity, and it became obvious that positive active materials having undergone complex substitution so as to comprise such compositions showed good cycle characteristics as a battery.

[0069] As described above, according to a lithium secondary battery of the present invention, sizable reduction in battery's internal resistance is realized since materials with improved electronic conductivity as well as low resistance which have been obtained with transition elements in a lithium transition metal compound having undergone complex substitution have been used as positive active materials. In addition, according to the present invention, reduction in positive capacity in excess of element substitution quantity is controlled. As a result of this, a lithium secondary battery according to the present invention serves to provide extremely excellent effects such as large output, huge capacity as

well as improved and good charge- discharge cycle characteristics, and further with less energy loss at the time of charging and discharging. Incidentally, in the case where LiMn_2O_4 has been used, such effect that reversibility of crystal configuration associated with charging and discharging is improved and superior endurance is provided is obtainable.

[0070] A lithium secondary battery has small internal resistance and has good charge-discharge cycle characteristics, with a lithium transition metal compound being used as a positive active material. A portion of transition element Me in a lithium transition metal compound LiMe_xO_y to be used as a positive active material is substituted by not less than two kinds selected from the group consisting of Li, Fe, Mn, Ni, Mg, Zn, B, Al, Co, Cr, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W. Here, M represents substitution elements, and $M \neq \text{Me}$.

10 **Claims**

1. A lithium secondary battery, comprising:

15 a lithium transition metal compound LiMe_xO_y , in which a portion of transition element Me is substituted by not less than two kinds selected from the group consisting of Li, Fe, Mn, Ni, Mg, Zn, B, Al, Co, Cr, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W to constitute $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ (herein M represents substitution elements, and $M \neq \text{Me}$, and Z represents quantity of substitution.), the $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ being to be used as a positive active material.

20 2. The lithium secondary battery according to claim 1, wherein the substitution elements M are not less than two kinds selected from the group consisting of Li, Fe, Mn, Ni, Mg, Zn, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W.

25 3. The lithium secondary battery according to claim 1 or 2, wherein at least Ti is included as the substitution elements M.

4. The lithium secondary battery according to claim 2 or 3, wherein a portion of the transition elements Me in $\text{LiM}_z\text{Me}_{x-z}\text{O}_y$ to include not less than two kinds of substitution elements M is substituted further by at least not less than one kind of elements selected from the group consisting of B, Al, Co, and Cr.

30 5. The lithium secondary battery according to any one of claims 1 to 4, wherein Z/X , a ratio of the substitution quantity Z and the transition element Me quantity X, fulfills a relationship of $0.005 \leq Z/X \leq 0.3$.

6. The lithium secondary battery according to any one of claims 1 to 5, wherein the lithium transition metal compound is lithium manganese oxide.

35 7. The lithium secondary battery according to claim 6, wherein an average valence of the substitution elements M to substitute a portion of manganese in the lithium manganese oxide is set at not less than 3 but not more than 4.

8. The lithium secondary battery according to claim 6 or 7, wherein the lithium manganese oxide has a spinel configuration of cubic system.

40 9. The lithium secondary battery according to any one of claims 6 to 8, wherein the substitution quantity Z remains within a range of $0.01 \leq Z \leq 0.5$.

10. The lithium secondary battery according to claim 9, wherein the substitution quantity Z remains within a range of $0.1 \leq Z \leq 0.3$.

45 11. The lithium secondary battery according to any one of claims 1 to 5, wherein the lithium transition metal compound is lithium-cobalt oxide or lithium nickel oxide.

50 12. The lithium secondary battery according to claim 11, wherein an average valence of the substitution elements M to be substituted with a portion of cobalt or nickel in lithium cobalt oxide or lithium nickel oxide is 3.

13. The lithium secondary battery according to claim 11 or 12, wherein the substitution quantity Z remains within a range of $0.005 \leq Z \leq 0.3$.

55 14. The lithium secondary battery according to claim 13, wherein the substitution quantity Z remains within a range of $0.05 \leq Z \leq 0.3$.

15. The lithium secondary battery according to any one of claims 1 to 14, wherein the $\text{LiM}_2\text{Me}_{x-z}\text{O}_y$ is composed by firing a mixed compound comprising salts and/or oxides having been prepared with a predetermined ratio in oxidation atmosphere at a range of 600°C to 1000°C, spending 5 hours to 50 hours.
- 5 16. The lithium secondary battery according to claim 15, wherein the $\text{LiM}_2\text{Me}_{x-z}\text{O}_y$ has been synthesized and obtained, conducting firing not less than twice, with the firing temperature for the forthcoming step to be set higher than that for the previous step.

10

15

20

25

30

35

40

45

50

55



(19)

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 0 973 217 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
28.06.2000 Bulletin 2000/26

(51) Int. Cl.⁷: H01M 4/48, H01M 4/50,
H01M 4/52, H01M 4/04

(43) Date of publication A2:
19.01.2000 Bulletin 2000/03

(21) Application number: 99113425.5

(22) Date of filing: 12.07.1999

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 13.07.1998 JP 19785398
20.11.1998 JP 33144998

(71) Applicant: NGK INSULATORS, LTD.
Nagoya-City, Aichi Prefecture 467-8530 (JP)

(72) Inventors:
• Nemoto, Hiroshi
Nagoya-city, Aichi-pref. 457-0022 (JP)
• Takahashi, Michio
Nagoya-city, Aichi-pref. 467-0059 (JP)
• Kitoh, Kenshin
Nagoya-city, Aichi-pref. 458-0827 (JP)

(74) Representative:
Bühling, Gerhard, Dipl.-Chem. et al
Patentanwaltsbüro
Tiedtke-Bühling-Kinne & Partner
Bavariaring 4
80336 München (DE)

(54) Lithium transition metal compound for lithium secondary battery

(57) A lithium secondary battery has small internal resistance and has good charge-discharge cycle characteristics, with a lithium transition metal compound being used as a positive active material. A portion of transition element Me in a lithium transition metal compound LiMe_xO_y to be used as a positive active material is substituted by not less than two kinds selected from the group consisting of Li, Fe, Mn, Ni, Mg, Zn, B, Al, Co, Cr, Si, Ti, Sn, P, V, Sb, Nb, Ta, Mo, and W to constitute $\text{LiM}_2\text{Me}_{x-z}\text{O}_y$. Here, M represents substitution elements, and M ≠ Me.

EP 0 973 217 A3



EUROPEAN SEARCH REPORT

Application Number
EP 99 11 3425

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	FR 2 738 673 A (MOLI ENERGY 1990 LTD) 14 March 1997 (1997-03-14) * page 5, line 15 - line 29 * * page 6, line 13 - line 21 * * page 15, line 5 - line 11 * * page 16, line 13 - line 20 *	1,2,6,8, 15	H01M4/48 H01M4/50 H01M4/52 H01M4/04
X	G. PISTOIA, A. ANTONUCCI, R. ROSATI ET AL: "Doped Li-Mn Spinels : Physical/Chemical characteristics and electrochemical performance in Li batteries" CHEMISTRY OF MATERIALS, vol. 9, no. 6, 1997, pages 1443-1450, XP002121765 * page 1443, left-hand column, paragraph 1 * * page 1443, right-hand column, paragraph 2 - paragraph 3 * * page 1444, left-hand column, paragraph 5 - paragraph 7; table 1 * * page 1445, left-hand column, paragraph 2 - page 1445, right-hand column, paragraph 1 * * page 1448, right-hand column, paragraph 2 * * table 7 *	1-6,8,9, 15	
-/-			
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	12 April 2000	Gamez, A	
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 99 11 3425

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	<p>PATENT ABSTRACTS OF JAPAN vol. 1998, no. 02, 30 January 1998 (1998-01-30) -& JP 09 270259 A (WAKIHARA MASATAKA), 14 October 1997 (1997-10-14) * abstract * <p>-& CHEMICAL ABSTRACTS, vol. 127, no. 22, 1 December 1997 (1997-12-01) Columbus, Ohio, US; abstract no. 309488, WAKIHARA, MASATAKA ET AL: "substituted spinel type lithium manganese oxide cathodes and secondary lithium batteries" XP002121766 * abstract * </p> </p>	1,5,8,9, 15	
X	<p>EP 0 782 206 A (JAPAN STORAGE BATTERY CO LTD) 2 July 1997 (1997-07-02) * page 2, line 45 - line 51 * * page 4, line 3 - line 13 * </p>	1,4,5, 11,13-15	
X	<p>EP 0 696 075 A (SEIKO INSTR INC) 7 February 1996 (1996-02-07) * page 3, line 10 - line 18 * * page 3, line 31 - line 41 * * page 5, line 35 - line 40 * * page 6, line 21 - line 29 * * page 6, line 51 - line 56 * * page 7, line 29 - line 38 * * page 8, line 29 - line 34 * * page 9, line 9 - line 14 * * page 10, line 44 - line 53 * * table 1 * </p>	1,2,4,5, 11,13-16	TECHNICAL FIELDS SEARCHED (Int.Cl.7)
<p>The present search report has been drawn up for all claims</p>			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	12 April 2000	Gamez, A	
CATEGORY OF CITED DOCUMENTS		<p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons -> : member of the same patent family, corresponding document</p>	
<p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p>			



DOCUMENTS CONSIDERED TO BE RELEVANT					
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.)		
X	WO 98 02928 A (ACCUMULATEURS FIXES) 22 January 1998 (1998-01-22) * page 2, line 1 - page 3, line 3 * * page 3, line 14 - line 25 * * page 4, line 23 - line 25 * * examples 6,7 *	1-3,5, 11,13-15			
X	EP 0 794 585 A (MITSUBISHI CABLE IND LTD) 10 September 1997 (1997-09-10) * page 3, line 17 - line 21 * * page 4, line 2 - line 6 * * table 2 *	1,11, 13-15			
X	PATENT ABSTRACTS OF JAPAN vol. 1995, no. 10, 30 November 1995 (1995-11-30) & JP 07 192719 A (SANYO ELECTRIC CO LTD), 28 July 1995 (1995-07-28) * abstract *	1,4,5, 11,13,14			
P, X	WO 98 38648 A (BARNETTE WAYNE ; FAULKNER TITUS (US); FMC CORP (US); GAO YUAN (US)) 3 September 1998 (1998-09-03) * page 3, line 13 - line 31 * * page 4, line 17 - line 23 * * page 5, line 23 - line 35 * * page 7, line 22 - page 8, line 11 * * page 11, line 27 - page 12, line 14 * * examples 1-3 *	1-6, 8-10,15, 16	TECHNICAL FIELDS SEARCHED (Int.Cl.)		
X	WO 98 06670 A (UEDA MASAMI ; FUKAMI TADASHI (JP); INOUE HIDETOSHI (JP); MATSUBARA) 19 February 1998 (1998-02-19) * page 3, paragraph 16 - paragraph 23 * * page 7, line 6 * * page 7, line 20 *	1,4,5, 11,13-15			
-/-					
The present search report has been drawn up for all claims					
Place of search	Date of completion of the search	Examiner			
THE HAGUE	12 April 2000	Gamez, A			
CATEGORY OF CITED DOCUMENTS					
X : particularly relevant if taken alone	T : theory or principle underlying the invention				
Y : particularly relevant if combined with another document of the same category	E : earlier patent document, but published on, or after the filing date				
A : technological background	D : document cited in the application				
O : non-written disclosure	L : document cited for other reasons				
F : intermediate document	& : member of the same patent family, corresponding document				



EUROPEAN SEARCH REPORT

Application Number

EP 99 11 3425

DOCUMENTS CONSIDERED TO BE RELEVANT					
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)		
X	EP 0 744 780 A (MATSUSHITA ELECTRIC IND CO LTD) 27 November 1996 (1996-11-27) * page 3, line 33 - line 35 * * page 4, line 18 - line 19 * * tables 9,20 *	1,4,5, 11,13-15			
X	WO 98 24131 A (POLYSTAR CORP) 4 June 1998 (1998-06-04) * page 4, line 29 - page 5, line 14 * * page 9, line 33 - page 10, line 1 * * page 10, line 14 - line 19 * * page 16, line 2 - line 10 * * example 4 *	1,3-5, 11,13-15			
P,X	YUAN GAO AND AL: "Novel LiNi _{1-x} Ti _x /2Mgx/2 compounds as cathode materials for safer lithium-ion batteries" ELECTROCHEMICAL AND SOLID-STATE LETTERS, vol. 1, no. 3, September 1998 (1998-09), pages 117-119, XP000859260 * page 117, left-hand column, paragraph 3 - paragraph 4 * * page 118, left-hand column, paragraph 3 * * page 119, right-hand column, paragraph 2 *	1-3,5, 11,13-15	TECHNICAL FIELDS SEARCHED (Int.Cl.7)		
X	PATENT ABSTRACTS OF JAPAN vol. 014, no. 028 (E-875), 19 January 1990 (1990-01-19) -& JP 01 265456 A (BRIDGESTONE CORP), 23 October 1989 (1989-10-23) * abstract *	1,2,5			
P,X	WO 98 57386 A (SANYO ELECTRIC CO ;NISHIO KOJI (JP); OHSHITA RYUKI (JP); KIDA YOSH) 17 December 1998 (1998-12-17) * tables 6,7 *	1-3,5, 11,13-15			
-/-					
The present search report has been drawn up for all claims					
Place of search	Date of completion of the search	Examiner			
THE HAGUE	12 April 2000	Gamez, A			
CATEGORY OF CITED DOCUMENTS					
X : particularly relevant if taken alone	T : theory or principle underlying the invention				
Y : particularly relevant if combined with another document of the same category	E : earlier patent document, but published on, or after the filing date				
A : technological background	D : document cited in the application				
O : non-written disclosure	L : document cited for other reasons				
P : intermediate document	& : member of the same patent family, corresponding document				



EUROPEAN SEARCH REPORT

Application Number

EP 99 11 3425

DOCUMENTS CONSIDERED TO BE RELEVANT									
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.)						
P, X	<p>WO 99 01903 A (FUJIWARA TAKAFUMI ; KAWANO TOMOKO (JP); HASHIMOTO AKIRA (JP); SYOJI) 14 January 1999 (1999-01-14)</p> <p>* page 5, line 9 - line 12 *</p> <p>* page 20, line 1 *</p>	1, 5, 11, 13-15							
TECHNICAL FIELDS SEARCHED (Int.Cl.)									
<p>The present search report has been drawn up for all claims</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 33%;">Place of search</td> <td style="width: 33%;">Date of completion of the search</td> <td style="width: 33%;">Examiner</td> </tr> <tr> <td>THE HAGUE</td> <td>12 April 2000</td> <td>Gomez, A</td> </tr> </table>				Place of search	Date of completion of the search	Examiner	THE HAGUE	12 April 2000	Gomez, A
Place of search	Date of completion of the search	Examiner							
THE HAGUE	12 April 2000	Gomez, A							
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>									

European Patent
Office

Application Number

EP 99 11 3425

CLAIMS INCURRING FEES

The present European patent application comprised at the time of filing more than ten claims.

Only part of the claims have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been paid, namely claim(s):

No claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims.

LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

see sheet B

All further search fees have been paid within the fixed time limit. The present European search report has been drawn up for all claims.

As all searchable claims could be searched without effort justifying an additional fee, the Search Division did not invite payment of any additional fee.

Only part of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the inventions in respect of which search fees have been paid, namely claims:

None of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the invention first mentioned in the claims, namely claims:



The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

1. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $\text{LiM}_{\text{x}}\text{O}_{\text{y}}$ in which a portion of transition element M_{x} is substituted by Li and at least one kind selected from the group consisting of Fe , Mn , Ni , Mg , Zn , B , Al , Co , Cr , Si , Ti , Sn , P , V , Sb , Nb , Ta , Mo and W to constitute $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-zO}_{\text{y}}$ (herein M represents Li and at least one other substitution element, and M different M_{x} , and z represents quantity of substitution), the $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-z}$ being to be used as a positive active material.

2. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $\text{LiM}_{\text{x}}\text{O}_{\text{y}}$ in which a portion of transition element M_{x} is substituted by at least one element of the triade VIIA of the periodic table selected from Fe , Ni and Co and at least one other element to constitute $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-zO}_{\text{y}}$, the $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-z}$ being to be used as a positive active material.

3. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $\text{LiM}_{\text{x}}\text{O}_{\text{y}}$ in which a portion of transition element M_{x} is substituted by at least one element of the group VIIA of the periodic table namely Mn and at least one other element to constitute $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-zO}_{\text{y}}$, the $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-z}$ being to be used as a positive active material.

4. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $\text{LiM}_{\text{x}}\text{O}_{\text{y}}$ in which a portion of transition element M_{x} is substituted by at least one element of the group IIA of the periodic table namely Mg and at least one other element to constitute $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-zO}_{\text{y}}$, the $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-z}$ being to be used as a positive active material.

5. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $\text{LiM}_{\text{x}}\text{O}_{\text{y}}$ in which a portion of transition element M_{x} is substituted by at least one element of the group IIB namely Zn and at least one other element to constitute $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-zO}_{\text{y}}$, the $\text{LiM}_{\text{z}}\text{M}_{\text{x}}\text{-z}$ being to be used as a positive active material.

LACK OF UNITY OF INVENTION
SHEET B

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

6. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $LiMexOy$ in which a portion of transition element Me is substituted by at least one element of the group IIIB selected from B and Al and at least one other element to constitute $LiMzMex-zOy$, the $LiMzMex-z$ being to be used as a positive active material.

7. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $LiMexOy$ in which a portion of transition element Me is substituted by at least one element of the group VIA selected from Cr, Mo and W and at least one other element to constitute $LiMzMex-zOy$, the $LiMzMex-z$ being to be used as a positive active material.

8. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $LiMexOy$ in which a portion of transition element Me is substituted by at least one element of the group IVB selected from Si and Sn and at least one other element to constitute $LiMzMex-zOy$, the $LiMzMex-z$ being to be used as a positive active material.

9. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $LiMexOy$ in which a portion of transition element Me is substituted by at least one element of the group IVA namely Ti and at least one other element to constitute $LiMzMex-zOy$, the $LiMzMex-z$ being to be used as a positive active material.

10. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $LiMexOy$ in which a portion of transition element Me is substituted by at least one element of the group VB selected from P and Sb and at least one other element to constitute $LiMzMex-zOy$, the $LiMzMex-z$ being to be used as a positive active material.

LACK OF UNITY OF INVENTION
SHEET B

Application Number

EP 99 11 3425

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

11. Claims: claim1-16(partially)

A lithium secondary battery, comprising : a lithium transition metal compound $LiMexOy$ in which a portion of transition element Me is substituted by at least one element of the group VA selected from V, Nb and Ta and at least one other element to constitute $LiMzMex-zOy$, the $LiMzMex-z$ being to be used as a positive active material.

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 99 11 3425

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

12-04-2000

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
FR 2738673	A	14-03-1997	CA 2158242 A DE 19636861 A GB 2305168 A, B JP 9147867 A US 5631104 A		14-03-1997 27-03-1997 02-04-1997 06-06-1997 20-05-1997
JP 09270259	A	14-10-1997	JP 2893327 B		17-05-1999
EP 0782206	A	02-07-1997	JP 9237631 A CN 1156910 A US 5718989 A US 5795558 A		09-09-1997 13-08-1997 17-02-1998 18-08-1998
EP 0696075	A	07-02-1996	JP 8213052 A DE 69514437 D US 5620812 A		20-08-1996 17-02-2000 15-04-1997
WO 9802928	A	22-01-1998	FR 2751135 A CA 2228671 A EP 0858677 A JP 11513181 T		16-01-1998 22-01-1998 19-08-1998 09-11-1999
EP 0794585	A	10-09-1997	JP 8153541 A AU 3936395 A US 5804335 A WO 9617392 A US 5989745 A		11-06-1996 19-06-1996 08-09-1998 06-06-1996 23-11-1999
JP 07192719	A	28-07-1995	NONE		
WO 9838648	A	03-09-1998	JP 2000500280 T		11-01-2000
WO 9806670	A	19-02-1998	CN 1232438 A EP 0918041 A		20-10-1999 26-05-1999
EP 0744780	A	27-11-1996	JP 9274917 A JP 9092285 A US 5631105 A		21-10-1997 04-04-1997 20-05-1997
WO 9824131	A	04-06-1998	US 5783333 A US 6007947 A		21-07-1998 28-12-1999
JP 01265456	A	23-10-1989	NONE		
WO 9857386	A	17-12-1998	EP 0989622 A		29-03-2000

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 99 11 3425

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

12-04-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9901903 A	14-01-1999	CN 1231067 T EP 0929111 A JP 11073966 A	06-10-1999 14-07-1999 16-03-1999

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82